



[10123/01101]

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant : Haarala et al.
Serial No. : 09/838,618
Filed : April 19, 2001
For : Catheter Slit Valves
Group Art Unit : 3754
Examiner : Buechner, Patrick M.

Mail Stop: Appeal Brief - Patent
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

APPEAL BRIEF UNDER 37 C.F.R. § 41.37

In support of the Notice of Appeal filed December 28, 2004, and pursuant to 37 C.F.R. § 41.37, Appellants present in triplicate their appeal brief in the above-captioned application.

This is an appeal to the Board of Patent Appeals and Interferences from the Examiner's final rejection of claims 43 - 46 and 61 in the final Office Action dated August 12, 2004 and the Advisory Action of November 18, 2004. The appealed claims are set forth in the attached Claims Appendix.

1. Real Party In Interest

This application is assigned to Scimed Life Systems, Inc., the real party in interest.

2. Related Appeals and Interferences

A Notice of Appeal and an Appeal Brief had been previously filed in response to the Office Action dated May 6, 2003. In view of the applicants' Appeal Brief, the Examiner reopened prosecution with new grounds for a non-final rejection prior to a decision on the merits by the Board of Patent Appeals and Interferences. *See*, the Office Action dated March 9, 2004. In response, the applicants elected to reply to the Examiner's rejection and did not request reinstatement of the prior appeal. Thus, the previous appeal has been vacated and will not have any bearing on the Board's decision in this appeal. The undersigned representative of the applicants is not aware of any other appeal or interference which will directly affect, or be directly affected by, or have a bearing on, the Board's decision in this appeal.

3. Status of Claims

Claims 1 - 42 and 47 - 60 were withdrawn from consideration. Claims 43 - 46 and 61 have been rejected in the final Office Action and are the subject of the instant appeal.

4. Status of Amendments

A Response to Final Rejection dated August 12, 2004 has been considered, but was not deemed to place the application in condition for allowance. *See*, the Advisory Action dated November 18, 2004.

5. Summary of Claimed Subject Matter

The invention is directed to a medical device comprising an elongate catheter including an external surface and at least one internal surface defining an internal lumen that extends longitudinally along at least a portion of the elongate catheter. See, Specification ¶ [0063]. A compound slit extends from a generally hemispherical portion of the external surface to the at least one internal surface and into communication with the internal lumen. See, Specification ¶ [0065]. The compound slit is biased toward a closed position and opens in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter. See, Specification, ¶ [0066].

6. Grounds of Rejection to be Reviewed on Appeal

- I. Whether claims 43, 44 and 61 are unpatentable under 35 U.S.C. § 102(b) as anticipated by Yoon (U.S. Patent No. 5,797,888).
- II. Whether claim 45 is unpatentable under 35 U.S.C. § 103(a) as obvious over Yoon in view of Phelps et al. (U.S. Patent No. 6,419,659).
- III. Whether claim 46 is unpatentable under 35 U.S.C. § 103(a) as obvious over Yoon in view of Desai (U.S. Patent No. 5,857,464).

7. Argument

- I. The Rejection of Claims 43, 44 and 61 Under 35 U.S.C. § 102(b) as Anticipated by Yoon (U.S. Patent No. 5,797,888) Should be Withdrawn.

It is respectfully submitted that applicants' medical device as recited in claim 43 is not anticipated by Yoon, as indicated by the Examiner. The Examiner stated, in support of the rejection that Yoon discloses all of the limitations of claim 43. Specifically, the Examiner asserted that Yoon disclosed a slit that is biased closed and would inherently open due to a difference in pressure between the lumen and the ambient. See, the Office Action dated August

12, 2004, 2. In addition, the Examiner alleged that Yoon is inherently configured to allow flaps to flex into the lumen when the ambient pressure exceeds the pressure inside the lumen. See id., 2.

Claim 43 recites a medical device comprising “an elongate catheter including an external surface and at least one internal surface defining an internal lumen that extends longitudinally along at least a portion of the elongate catheter” and “a compound slit extending from a generally hemispherical portion of the external surface to the at least one internal surface and into communication with the internal lumen, the compound slit being biased toward a closed position and opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter.”

The Examiner has asserted that the slit of Yoon would inherently open due to differences in fluid pressure between the lumen and the ambient. However, “[i]n relying upon the theory of inherency, the examiner must provide a basis in fact and/or technical reasoning to reasonably support the determination that the allegedly inherent characteristic necessarily flows from the teachings of the applied prior art.” Ex parte Levy, 17 USPQ2d 1461, 1464(Bd. Pat. App. & Inter. 1999). The Examiner has asserted that the above-identified characteristic is inherent, but neglected to provide a basis in fact or a technical reasoning to reasonably support her statement. Therefore, it is respectfully submitted that the Examiner has not met the requirements for relying upon the theory of inherency and thus, has not provided sufficient basis for her § 102(b) rejection.

Additionally, the Examiner is directed to the requirements for establishing inherency.

“To establish inherency, the extrinsic evidence ‘must make clear that the missing descriptive matter is necessarily present in the thing described in the reference and that it would be so recognized by persons of ordinary skill. Inherency, however, may not be established by probabilities or possibilities. The mere fact that a certain thing may result from a given set of circumstances is not sufficient.’” In re Robertson, 169 F.3d 743, 745 (Fed. Cir. 1999)(citations omitted).

It is also respectfully submitted that Yoon does not make clear that the slit would necessarily open due to differences in the fluid pressure of the lumen and fluid pressure outside the catheter.

Specifically, Yoon describes a cannula which includes a seal, "normally biased to a closed position when no instrument is passed through the cannula...." Yoon Patent, col. 4, lines 44-46. The only means discussed in Yoon's disclosure for opening the seal is by pushing a medical instrument or a pusher through the seal from the interior of the cannula in order to overcome the closing bias of the seal. See id., col. 6, lines 57-62. The seal opens to accommodate and substantially conform to the shape of the instrument, but returns to a closed position upon removal of the instrument from the seal. See id., col. 4, lines 58-62. Similarly, the seal also opens to substantially conform to the exterior of a tubular pusher. See id., col. 7, lines 4-9. However, the seal is biased to a closed position and "once the pusher has returned to the retracted position, the seal will no longer be held in an open position and will be free to move to a sealing position with seal members urged toward the closed position...." Id., col. 7, lines 31-34 (citations omitted). Yoon makes no reference to a compound slit that opens in response to the difference between a fluid pressure within the lumen and a fluid pressure outside the catheter. Rather, as discussed above, the Yoon reference makes it clear that the seal is to remain closed at all times when no pusher or instrument is pushed therethrough to force the sealing members to an open position. Opening a slit in response to a difference of fluid pressures between the interior and the exterior of the catheter is unrelated to and does not necessarily flow from opening a seal by overcoming the closing bias with a pusher or a medical instrument. Specifically, at the pressures to which such medical devices are exposed, this seal is designed to remain closed. Of course the seal eventually would open if the pressure applied thereto were continually increased. However, the solid walls of the catheter would also eventually open under these conditions. It is respectfully submitted that such strained interpretations are not indicative of the understanding which would be developed by a person skilled in the art upon reading Yoon. Thus, appellants respectfully submit that there is no inherent disclosure of a compound slit opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter.

The present invention as disclosed by claim 43 refers to a medical device including a compound slit biased toward a closed position and opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter. The compound slit of the appellants' invention opens and permits fluid flow through the catheter in response to an increased pressure differential between the lumen and the external environment. For example, as described in the specification, in an infusion mode, "the pressure within the lumen 228 is increased with respect to the environmental pressure, thereby causing the catheter walls 227 adjacent the slit 222 to begin to move apart, allowing infusion of fluid to the external environment." Specification, ¶[0078]. Furthermore, the catheter may alternatively be configured to an aspiration mode, where "[a]s the lumen pressure decrease[s] with respect to the environmental pressure, one side of the slit valve 222 yields, or flexes inwardly, to allow fluid to enter the catheter lumen 228." Id., ¶[0078]. As would be understood by one of ordinary skill in the art, the slit described by the appellants' application may open to permit fluid flow without placing a physical structure (i.e., a pusher or a medical instrument) through the slit to overcome the closing bias of the sealing members.

Applicants respectfully submit that the Examiner's reading of Yoon is directly contradictory to the teachings contained therein. In particular, Yoon claims that fluid flow between the interior of the cannula and the exterior environment is undesirable and prohibited by the seal. See Yoon Patent, col. 10, lines 3-6. Yoon makes clear that the seal prevents fluid flow through the cannula; particularly when the sealing members are not forced to an open configuration by the insertion of a pusher or a medical instrument. See id., col. 10, lines 10-14. More specifically, "[t]he seal can have any configuration to prevent fluid flow through the cannula prior to the introduction of instruments through the cannula, after the instruments are withdrawn from the cannula and/or while the instruments are in place." Id., col. 9, lines 53-57. One of ordinary skill in the art would understand that fluids generally flow from an area of high concentration / pressure to an area of low concentration / pressure. Accordingly, because Yoon's seal is designed to prevent fluid flow across the seal, the seal must serve as a barrier between two areas of different fluid pressures. Therefore, appellants respectfully submit that Yoon does not

disclose a seal that succumbs to fluid pressure gradients and opens in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter.

The Yoon reference further illustrates that the seal is not equivalent to the compound slit which opens in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter, as recited by claim 43 of the applicants' invention. Specifically, Yoon describes the use of a plurality of spring wires or stiffeners to maintain the seal in a closed position. See *id.*, col. 4, lines 62-67 and col. 5, lines 1-4. Use of the spring wires in the seal would prevent anything other than a pusher or a medical instrument from overcoming the closing bias and forcing open the seal. It is respectfully submitted that there is no disclosure that would lead one to conclude that the resilient force of the spring wires and other structures biasing the slits closed would open under the influence of any fluid pressure gradients as are encountered in the anatomical environments to which medical devices, such as the present invention, are exposed.

In its specification, Yoon discussed the use of a distensible or inflatable membrane as disclosed in U.S. Patent Application Serial No. 09/383,520 ('520 Application), which is now abandoned. See Yoon Patent, col. 11, lines 4-6. Furthermore, Yoon incorporated the disclosure of the abandoned '520 application by reference. See *id.*, col. 11, lines 5-8. U.S. Patent No. 5,752,970 to Yoon ('970 Patent) is a continuation thereof and thus, shares the same disclosure as the '520 application. In the '970 patent, an expandable membrane, the same as that incorporated by Yoon, is used to seal the distal end. Thus, Yoon includes the membrane as an element of the seal. Therefore, characterizations of the seal disclosed by the '970 patent apply to Yoon as well. The disclosure of the '970 patent further brings to light that the seal recited by Yoon does not permit opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter. In fact, the disclosure of the '970 patent expressly states to the contrary that "[t]he valves are preferably configured as one-way valves such that external forces and pressures exerted on the valves from outside the cannula will not cause the valves to open." '970 Patent, col. 7, lines 59-62 (emphasis added). Therefore, in

view of the disclosure incorporated by reference, Yoon clearly teaches away from a compound slit opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter.

The Examiner asserts that “[t]he slit of Yoon is also configured to inherently allow the flaps to flex into the lumen when the ambient pressure exceeds the pressure inside the lumen.” Appellants respectfully disagree with the Examiner’s statement. The seal in Yoon never opens in response to an inward force and only opens outwardly a result of being physically pushed by the pusher or the instrument. In addition, the seal acts to prevent all fluid flow back into the cannula. One of ordinary skill in the art would ascertain that the pusher or instrument exerts only an outward physical force against the seal from the interior of the cannula and that the seal does not open in response to inward forces, such as fluid flow back into the cannula. Consequently, it is respectfully submitted that that the device of Yoon teaches away from the configuration claimed by the applicants. Appellants further submit that Yoon does not teach a slit that opens in response to a pressure gradient, because such a slit would respond to both inward and outward forces.

The appellants have shown above that Yoon does not contemplate opening the seal in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter. In fact, Yoon has repeatedly taught away from a seal that responds to change in fluid pressure. In addition, Yoon has also taught away from the configuration of the seal as claimed by the applicants. Accordingly, it is respectfully submitted that Yoon does not make clear that the description of a slit that is biased closed and opens due to a difference in pressure between the lumen and the ambient as well as a slit configured to allow flaps to flex into the lumen when the ambient pressure exceeds the pressure inside the lumen are necessarily present in Yoon’s disclosure and that it would be so recognized by persons of ordinary skill. Thus, the appellants respectfully submit that these elements are not inherent within the Yoon reference.

"A claim is anticipated only if each and every element as set forth in the claim is found, either expressly or inherently described, in a single prior art reference." Verdegaal Bros. v. Union Oil Co. Of California, 814 F.2d 628, 631 (Fed. Cir. 1987). It is respectfully submitted that Yoon does not show or suggest each and every element of a medical device comprising an elongate catheter including an external surface and at least one internal surface defining an internal lumen that extends longitudinally along at least a portion of the elongate catheter; and a compound slit extending from a generally hemispherical portion of the external surface to the at least one internal surface and into communication with the internal lumen, the compound slit being biased toward a closed position and opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter as recited in amended claim 43.

Accordingly, applicants respectfully submit that Yoon does not disclose, either expressly or inherently, each and every element of independent claim 43 and that this rejection should be withdrawn. Because claims 44 and 61 depend from and, therefore, include all of the limitations of claim 43, it is respectfully submitted that these claims are also allowable for the same reasons as indicated above.

II. The Rejection of Claim 45 Under 35 U.S.C. § 103(a) as Unpatentable Over Yoon in View of Phelps (U.S. Patent No. 6,419,659) Should be Withdrawn.

It is respectfully submitted that the appellants' medical device as recited in claim 45, is not obvious over Yoon in view of Phelps et al. for substantially the same reasons stated above in regard to claims 43, 44 and 61.

The Examiner stated, in support of the rejection that Yoon as described above shows a device substantially as claimed except for the element of a collar disposed adjacent to the distal end of the catheter, but that Phelps et al. discloses a collar disposed adjacent the catheter's distal-most end. The Examiner further stated that it would have been obvious for one

of ordinary skill in the art to combine the above-mentioned prior art and that “[d]oing so would provide an attending physician with a means for determining the location of the catheter by magnetic or electromagnetic means.” *See* Office Action dated August 12, 2004, 3 (citations omitted).

It is respectfully submitted that claim 45 is allowable for at least the same reasons stated above in regard to claims 43, 44 and 61, and that Phelps et al. fails to cure the noted defects.

III. The Rejection of Claim 45 Under 35 U.S.C. § 103(a) as Unpatentable over Yoon in view of Desai (U.S. Patent No. 5,857,464) Should be Withdrawn.

It is respectfully submitted that the appellants’ medical device as recited in claim 46, is not obvious over Yoon in view of Desai for substantially the same reasons stated above in regard to claims 43, 44 and 61.

The Examiner stated, in support of the rejection that Yoon as described above shows a device substantially as claimed except for the element of a tricuspid flap configuration, but Desai discloses a valve with three flaps. The Examiner further stated that it would have been obvious for one of ordinary skill in the art to combine the above-mentioned prior art and that “it would be a simple matter of choosing a design for an end valve from existing designs known in the art, when each design would perform equally well.” *See* Office Action dated August 12, 2004, 3.

In addition, applicants respectfully submit that neither of the cited references provides any suggestion, incentive or motivation for the combination as suggested by the Examiner. “Multiple cited prior art references must suggest the desirability of being combined and the reference must be viewed without the benefit of hindsight afforded to the disclosure.”(emphasis added) In re Paulsen, 30 F.3d 1475, 1482 (Fed. Cir. 1994). Applicants respectfully submit that *there is no motivation or incentive to combine* Yoon with Desai to

allegedly teach or suggest Applicants' claimed invention.

As stated by the Federal Circuit, "the teaching or suggestion to make the claimed combination and the reasonable expectation of success must both be found in the prior art, and not based on Applicant's disclosure." See In re Vaeck, 947 F.2d 488 (Fed. Cir. 1991).

Yoon teaches away from the proposed combination. Desai discloses a valve designed to open when acted upon by the force of fluid pressure to permit the guidewire to pass therethrough. Contrary to Desai, as discussed above, Yoon's seal opens only when a pusher or an instrument is physically pushed against the seal and overcoming the closing bias. The disclosure of Yoon makes it clear that seal serves to prevent fluid flow and will not succumb to fluid pressure.

As described above, Yoon prohibits fluid flow through the catheter even when opened by a pusher or instrument extending through the sealing members by tightly sealing around the pusher or instrument. Withdrawing the guidewires will seal the valve at the distal end of Desai's catheter as well. However, contrary to Yoon, Desai continues to evacuate liquids from the interior of the catheter to a cavity formed by intermediate portions of a slitted catheter tube, even subsequent to closing the distal valve. See Desai Patent, col. 5, lines 55-58. As described above, Desai does not restrict fluid flow in the same manner. Thus, since Yoon and Desai describe contradictory devices, it is respectfully submitted that neither of these references provides any suggestion, incentive or motivation for the combination and that this combination is an improper hindsight reconstruction.

It is respectfully submitted that neither Yoon nor Desai shows or suggests a medical device comprising an elongate catheter including an external surface and at least one internal surface defining an internal lumen that extends longitudinally along at least a portion of the elongate catheter; and a compound slit extending from a generally hemispherical portion of the external surface to the at least one internal surface, wherein the compound slit is a tricuspid

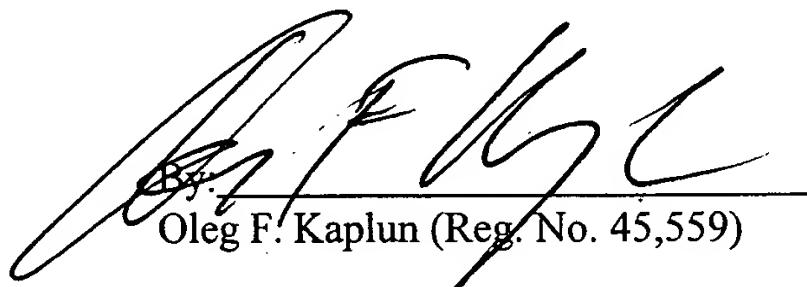
slit as recited in claim 46 .

Applicants respectfully submitted that claim 46 is allowable for at least the same reasons stated above in regard to claims 43, 44 and 61, and that Desai fails to cure the noted defects. Furthermore, it is respectfully submitted that claim 46 is neither shown nor suggested by the above-cited references either taken alone or in combination and that this rejection should be withdrawn.

8. Conclusion

It is respectfully submitted that appellants have demonstrated that the subject matter of independent claim 43 and those dependent therefrom (claims 44 and 61) are not anticipated by the cited prior art. In addition, appellants further submit that claims 45 and 46 are not obvious in light of the cited art, taken alone or in combination. Thus, it is respectfully requested that the Examiner's final rejection of claims 43 - 46 and 61 be reversed and all appealed claims found patentable.

Respectfully submitted,



Oleg F. Kaplun (Reg. No. 45,559)

Date: September 13, 2005

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CLAIMS APPENDIX

43. A medical device comprising:
 - an elongate catheter including an external surface and at least one internal surface defining an internal lumen that extends longitudinally along at least a portion of the elongate catheter; and
 - a compound slit extending from a generally hemispherical portion of the external surface to the at least one internal surface and into communication with the internal lumen, the compound slit being biased toward a closed position and opening in response to a difference between a fluid pressure within the lumen and a fluid pressure outside the catheter.
44. A medical device according to claim 43, wherein the compound slit is disposed on a distal end of the elongate catheter.
45. A medical device according to claim 44, further comprising a collar disposed at the distal end of the catheter.
46. A medical device according to claim 43, wherein the compound slit is a tricuspid slit.
61. A medical device according to claim 43, wherein the compound slit is configured so that, when the fluid pressure within the lumen exceeds the fluid pressure outside the catheter by a first predetermined amount, flaps of the hemispherical portion formed by the compound slit flex outward away from a longitudinal axis of the catheter to allow fluid within the lumen to exit and when the fluid pressure outside the catheter exceeds the fluid pressure within the lumen by a second predetermined amount, the flaps flex into the lumen to allow fluid outside the catheter to enter the lumen.

EVIDENCE APPENDIX

No evidence has been entered or relied upon in the present appeal.

RELATED PROCEEDING APPENDIX

No decisions have been rendered regarding the present appeal or any proceedings related thereto.



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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/838,618	04/19/2001	Brett T. Haarala	10123/01101	3578
7590	09/09/2005		EXAMINER	
Fay Kaplun & Marcin, LLP 150 Broadway Suite 702 New York, NY 10038			BUECHNER, PATRICK M	
			ART UNIT	PAPER NUMBER
			3754	

DATE MAILED: 09/09/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

**Notification of Non-Compliant Appeal Brief
(37 CFR 41.37)**

Application No.	09/838,618	Applicant(s)	HAARALA ET AL.
Examiner	Eric Keasel	Art Unit	3754

--The MAILING DATE of this communication appears on the cover sheet with the correspondence address--

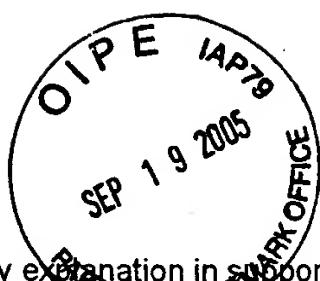
The Appeal Brief filed on 6/2 May 2005 is defective for failure to comply with one or more provisions of 37 CFR 41.37



1. The brief does not contain the items required under 37 CFR 41.37(c), or the items are not under the proper heading or in the proper order.
2. The brief does not contain a statement of the status of all claims, (e.g., rejected, allowed, withdrawn, objected to, canceled), or does not identify the appealed claims (37 CFR 41.37(c)(1)(iii)).
3. At least one amendment has been filed subsequent to the final rejection, and the brief does not contain a statement of the status of each such amendment (37 CFR 41.37(c)(1)(iv)).
4. (a) The brief does not contain a concise explanation of the subject matter defined in each of the independent claims involved in the appeal, referring to the specification by page and line number and to the drawings, if any, by reference characters; and/or (b) the brief fails to: (1) identify, for each independent claim involved in the appeal and for each dependent claim argued separately, every means plus function and step plus function under 35 U.S.C. 112, sixth paragraph, and/or (2) set forth the structure, material, or acts described in the specification as corresponding to each claimed function with reference to the specification by page and line number, and to the drawings, if any, by reference characters (37 CFR 41.37(c)(1)(v)).
5. The brief does not contain a concise statement of each ground of rejection presented for review (37 CFR 41.37(c)(1)(vi))
6. The brief does not present an argument under a separate heading for each ground of rejection on appeal (37 CFR 41.37(c)(1)(vii)).
7. The brief does not contain a correct copy of the appealed claims as an appendix thereto (37 CFR 41.37(c)(1)(viii)).
8. The brief does not contain copies of the evidence submitted under 37 CFR 1.130, 1.131, or 1.132 or of any other evidence entered by the examiner and relied upon by appellant in the appeal, along with a statement setting forth where in the record that evidence was entered by the examiner, as an appendix thereto (37 CFR 41.37(c)(1)(ix)).
9. The brief does not contain copies of the decisions rendered by a court or the Board in the proceeding identified in the Related Appeals and Interferences section of the brief as an appendix thereto (37 CFR 41.37(c)(1)(x)).
10. Other (including any explanation in support of the above items):

See Continuation Sheet.

Eric Keasel 6 SEP 2005
Eric Keasel
Primary Examiner
Art Unit: 3754



Continuation of 10. Other (including any explanation in support of the above items):

1. The new rules governing appeals (37 CFR 1.136) became effective last year and are required for all appeal briefs filed after September 2004. Appellant's representative should obtain a copy of the current rules.
2. The appeal brief filed on May 2, 2005 is not fully responsive to the prior Notification of Non-Compliant Appeal Brief because appellant has not complied with the requirements set forth in 37 CFR 41.37. Since the period for reply set forth in the prior Notification of Non-Compliant Appeal Brief has expired, this application will become abandoned unless applicant corrects the deficiency and obtains an extension of time under 37 CFR 1.136(a). The date on which the petition under 37 CFR 1.136(a) and the appropriate extension fee have been filed is the date for purposes of determining the period of extension and the corresponding amount of the fee. In no case may an applicant reply outside the SIX (6) MONTH statutory period or obtain an extension for more than FIVE (5) MONTHS beyond the date for reply set forth in the prior Notification of Non-Compliant Appeal Brief. A fully responsive reply must be timely filed to avoid abandonment of this application.
3. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Eric Keasel whose telephone number is (571) 272-4929. The examiner can normally be reached on Monday-Friday. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Mar can be reached on (571) 272-4906. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300. Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).